OMB Comment: OMB is required to make a decision concerning the collection of information between 30 to 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, NW., Attn: ACF Desk Officer.

Dated: August 18, 1999.

Bob Sargis,

Acting Reports Clearance Officer. [FR Doc. 99–218860 Filed 8–23–99; 8:45 am] BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Adoption and Foster Care Analysis and Reporting System for Title IV–B and Title IV–E.

OMB No.: 0980-0267.

Description: Section 479 of title IV-E of the Social Security Act directs States to establish and implement an adoption and foster care reporting system. The purpose of the data collected is to inform State/Federal policy decisions, program management, and to respond to

Congressional and Department inquiries. Specifically, the data is used for short/long-term budget projections, trend analysis, and to target areas for improved technical assistance. The data will provide information about foster care placements, adoptive parents, length of time in care, delays in termination of parental rights and placement for adoption.

Respondents: State, Local or Tribal Govt.

Annual Burden Estimates:

Instrument	Number of respondents		Average burden hours per response	Total bur- den hours
Adoption and Foster Care Analysis and Reporting Systems	51	2	3,251	331,.602

Estimated Total Annual Burden Hours: 331,602.

Additional Information: Copies of the proposed collection may be obtained by writing to The Administration for Children and Families, Office of Information Services, Division of Information Resource Management Services, 370 L'Enfant Promenade, SW; Washington, DC 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 to 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect is OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, NW., Washington, D.C. 20503, Attn: ACF Desk Officer.

Dated: August 18, 1999.

Bob Sargis,

Acting Reports Clearance Officer.
[FR Doc. 99–21894 Filed 8–23–99; 8:45 am]
BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N-1392]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; State Enforcement Notification

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments on the collection of information by September 23, 1999.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Wendy Taylor, Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

State Enforcement Notification—21 CFR 100.2(d) (OMB Control Number 0910-0275)—Extension

Section 310(b) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 337(b)) authorizes States to enforce certain sections of the act in their own names, but provides that States must notify FDA before doing so. Section 100.2(d) (21 CFR 100.2(d)) sets forth the information that a State must provide to FDA in a letter of notification when it intends to take enforcement action under the act against a particular food located in the State. The information required under § 100.2(d) will enable FDA to identify the food against which the State intends to take action and advise the State whether Federal action has been taken against it. With certain narrow exceptions, Federal enforcement action precludes State action under the act.

In the **Federal Register** of June 8, 1999 (64 FR 30525), the agency requested comments on the proposed collections of information. No comments were received.

FDA estimates the burden of this collection of information as follows: